California Medical Device Recall Information

Recall Name
Bard Peripheral Vascular Recalls Bard LifeStent Solo Vascular Stent Due to Deployment Failure

Recall Date	Product Description	Recalling Firm	Recall Reason
09/30/13	Bard LifeStent Solo Vascular Stent	**Bard Peripheral Vascular, Inc.** Tempe, AZ	*The deployment mechanism may not perform properly including: failure to deploy, partial deployment, and difficult deployment.* *This may result in adverse health consequences.*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
I	Item Numbers: • **EX062001CL** • **EX072001CL** • **EX062003CL** • **EX072003CL** For Affected Lots Click Here	Nationwide	Manufactured and distributed from: November 2011 to June 30, 2012.

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm371318.htm